

D'YAKONOV, I.A.; GOLCDNIKOV, G.V.; REPINSKAYA, I.B.

Reaction of diazoacetic ester with trimethylvinylsilane  
catalyzed by copper sulfate. Zhur.ob.khim. 32 no.10:3450-3451  
O '62. (MIRA 15:11)

1. Leningradskiy gosudarstvennyy universitet.  
(Serine) (Silane)

GRIGORYEVICH, V. I.

MA Biological Institute for the Study of the Field of Microbiology, 1961, 1962, 1963,  
Inst of Agriculture, Acad Sci USSR, Tashkent, 1961. 1962, 1963, 1964

1961: 1962, 1963, 1964

BORISOVA, A.G.; BOCHANTSEV, V.P.; BUTKOV, A.Ya., dotsent; VASIL'KOVSKAYA, A.P.;  
 VVEDENSKIY, A.I., dotsent; GOLODKOVSKIY, V.L.; GONCHAROV, N.F.  
 [deceased]; DROBOV, V.P., professor; KOROTKOVA, Ye.Ye.; KOSTINA, K.F.;  
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 MIRONOV, B.A. [deceased]; PAZIY, V.K.; POYARKOVA, A.I.; PROTOPOPOV,  
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 KOROVIN, Ye.P., professor, glavnyy redaktor; ZAKIROV, K.Z., professor,  
 redaktor; SHIPUKHIN, A.Ya, redaktor izdatel'stva

[The flora of Uzbekistan] Flora Uzbekistana. Glav. red. Ye.P.Korovin.  
 Tashkent, Izd-vo Akademii nauk UzSSR. Vol.3. 1955. 825 p. (MLA 9:10)

1. Deystvitel'nyy chlen AN UzSSR (for Korovin)  
 (Uzbekistan---Botany)

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Determining irrigation requirements of alfalfa grown for seed  
production. Dokl. AN Uz. SSR no. 9: 39-41 '56. (MIRA 12:6)

1. Institut sel'skogo khozyaystva AN UzSSR. Predstavleno chlenom-  
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(Alfalfa--Water requirements)  
(Seed production)

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GOLODKOVSKIY, V.L.; SHAAKRAMOV, K.Sh.

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1. Institut genetiki i fiziologii rasteniy AN UzSSR.  
(GRASSES)

GOLODKOVSKIY, V.L.; POPOVA, Ye.A.

In memory of V.S.Shardakov. Uzb. biol. zhur. no.1:77-78 '61.  
(MIRA 14:3)

(SHARDAKOV, VASILII SEMENOVICH, 1901-1960)

GOLOVCHENKO, S.G.; GOLODKOVSKIY, V.L., kand.sel'khoz.nauk, otv. red.;  
GOR'KOVY, P.I., red.; KARABAYEVA, Kh.U., tekhn. red.

[Feed production in piedmont areas; research results] Kernodoby-  
vanie v predgor'iax; opyt issledovaniia. Tashkent, Izd-vo Akad.  
nauk UzSSR, 1962. 166 p. (MIRA 15:7)  
(Uzbekistan--Forage plants)



**"APPROVED FOR RELEASE: 09/24/2001**

**CIA-RDP86-00513R000515730004-6**

**APPROVED FOR RELEASE: 09/24/2001**

**CIA-RDP86-00513R000515730004-6"**

GOLEKOVSKIY, V.I.; PATERIKHAYEV, S.I.

Heterosis in crossing self-pollinated lines of corn. Dokl.  
biol. zhur. 9 no.1:65-66 1965. (MIA 151c)

L. Institut eksperimental'noy biologii tekhnicheskikh i selskoykh  
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PALEY, M.L., INDIANA, L.A., IN, LONDON, L.A., COLOMBIA, S.L.

Investigation of resistance to fungi of the acoustical  
material based on polyvinyl chloride resin. First, massy  
no. 1000-1000. (MIRA 1974)

PELIN, I.M.; PLECHINSKY, V.I.; GILBERT, A.L.; PERCHINSKY, A.L.;  
KOTCHENKO, A.L.

Removal of Carcinogenic from the waste water of the production  
of carbanilic acid. Inst. org. n. 155-156, 1971, 155-156, 155-156, 155-156.



GOLODNIKOV, A. V.

Academy/Univ articles - economic

100-50

Annual scientific section of Leningrad University, in 1961, A. V. Golodnikov,  
T. A. Kozlov, A. G. Kozlov, A. V. Kozlov, and A. V. Kozlov.

Vest Leningrad U, Ser Nat, Soc, Econ, Vol 7, No 2, pp 188-190

The annual scientific section of Leningrad University was held on 9-10 Feb 1961.  
The Math section was subdivided into math, mechanics, and astronomy; the physics  
comprised also applied physics. The chemistry section dealt also with cooperation with  
industry.

LA 2517 6

*GOLODNIKOV, G.V.*

USSR/Chemistry - Catalysis

Card 1/1    Pub. 151 - 12/33

Authors    :   Dolgov, B. N., and Golodnikov, G. V.

Title       :   Catalytic derivation of mixed ketones from primary alcohols. Part 1.-  
Ketonization of an ethyl and n-butyl alcohol mixture

Periodical :   Zhur. ob. khim. 24/6, 987-993, June 1954

Abstract   :   Experiments on the ketonization of an ethyl and n-butyl alcohol mixture, carried out in the presence of additionally introduced hydrogen in molar ratio of 1 : 1 to the alcohol mixture, are described. The effect of the hydrogen addition on the ketone yield and catalyst activity is explained. The conditions favorable for such ketonization process and the experimental results obtained are listed. Nine references: 6-USSR and 3-USA (1924-1951). Tables.

Institution :   The A. A. Zhdanov State University, Leningrad

Submitted   :   December 26, 1953

GOLODNIKOV, G. V.

3  
CII Catalytic preparation of mixed ketones from primary  
alcohols. II. Ketoneization of a mixture of ethyl and iso-  
amyl alcohols. B. N. Dalgov and G. V. Golodnikov.  
J. Gen. Chem. U.S.S.R. 24, 1169-61 (1954) (Engl. Transla-  
tion).—See C.A. 49, 12282a. B. M. R. (1) Ph



Golodnikov, G. V.

USSR .

62  
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/ Catalytic preparation of mixed ketones from primary  
alcohols. II. Ketoneization of a mixture of ethyl and isopropyl  
alcohols. B. N. Dolgov and G. V. Golodnikov (Leningrad  
State Univ.). *Zhur. Obshch. Khim.* 24, 1107-70 (1954);  
cf. C.A. 49, 8794f.—Passage of EtOH and iso-AmOH in  
2:1 molar ratio over Cu catalyst activated according to  
directions given by D. and Bolotov (*Vestnik Leningrad.  
Gosudarst. Univ.* 5, No. 6 (1950) at 350° and space velocity  
50-150 gives the best yield (18-21%) of iso-BuAc, with 0.6-  
1% Me<sub>2</sub>CO and about 5% (iso-Bu)<sub>2</sub>CO. At 375-300°  
are formed esters in yields near 45%; these include EtOAc,  
iso-AmOAc, iso-BuCO<sub>2</sub>Et and iso-BuCO<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>Me.  
The effluent gases from the ketoneization contain mostly H<sub>2</sub>,  
with small amounts of CO, CO<sub>2</sub> and olefins. G. M. K.

G. V. Golodnikov, G. V.

USSR

Catalytic preparation of mixed ketones from primary alcohols. III. The ketonization of a mixture of *n*-BuOH and iso-AmOH. B. N. Dolgov and G. V. Golodnikov (Leningrad State Univ.). *Zh. Obshch. Khim.* 24, 1861-71 (1954); cf. *Chem. Abstr.* 49, 5947. In the passage of mixed *n*-BuOH and iso-AmOH over activated Cu catalyst at 275-425°, the best yields (18%) of  $\text{PrCOCH}_2\text{CHMe}_2$ , 23%  $\text{Pr}_2\text{CO}$ , and 2-3% iso-Bu<sub>2</sub>CO are obtained when the molar ratio of *n*-BuOH to iso-AmOH is 2-3:1 and the temp. is 350-75°, with a space velocity of 100-50. At lower temps. (275-325°) the main products are esters forming up to 60% of total product. The off-gas consists mainly of H<sub>2</sub> with small amts. of CO, CO<sub>2</sub>, and olefins. The results indicate that aldehydes and esters are the intermediate products of ketonization. The correct mechanism appears to be that suggested by Sabatier-Kagan (no ref. cited), in which esters are intermediates. The reaction yields aldehydes, which condense to esters which are then decarboxylated with elimination also of ROH and olefins, along with the H<sub>2</sub> formed in the initial dehydrogenation of ROH. G. M. Kosolapov

CH  
①

MA  
Met

GOLODNIKOV, G. V.

7  
 ✓ Decomposition of esters of benzoic acid over chromium catalyst. B. M. Dolgov, G. V. Golodnikov, and L. M. Chernykhina (State Univ., Leningrad, USSR, *Dokl. Akad. Nauk SSSR*, 1955, 60 (1955)).—Passage of  $\text{BzOPh}$ ,  $\text{BzOBu}$ , and  $\text{BzOAm}$  over  $\text{Zn-Cr}$  catalyst (cf. C.I. 49, 10170c) at  $330-420^\circ$  yields  $\text{C}_6\text{H}_6$ ,  $\text{CO}$ , and olefin, in one reaction sequence, and an alkyl aryl ketone in another sequence. For prepn. of the ketones the best conditions are  $100-200^\circ$  at space velocity 50 G; under these conditions the yields of ketones are:  $\text{AcPh}$  0.3%,  $\text{PhCOPh}$  6.6%,  $\text{iso-BuCOPh}$  2.3%.  
 G. M. Kosolapoff

USSR/Organic Chemistry. Synthetic Organic Chemistry.

E-2

Abs Jour: Ref Zhur-Khimiya, No 6, 1957, 19254

Author : Dolgov N. N., Golodnikov G. V., Glushkova N. Ye.

Inst : *Academy of Sciences of the USSR*

Title : Conversion of Tetraethylsilane over a Chromium Catalyst.

Orig Pub: Zh. obshch. khimii, 1956, 26, No 6, 1688-1691

Abstract: At 530-600° tetraethylsilane (I) over a chromium catalyst in an atmosphere of  $H_2$  is decomposed into triethylsilane and  $C_2H_4$ . At 600-630° a deep hydrogenolysis ensues with the formation of  $C_2H_5$  and Si. Dehydration of I to triethylvinylsilane is not observed.

Card : 1/1

GOLODNIKOV, G. V.

✓ Transformation of International Law  
B. S. Dolzov, G. V. Golodnikov, and M. L. Golodnikov  
J. Gen. Chem. 1955, 20, 1111-5 (1955) (English trans-  
lation) - See C. I. 11 1111

1111

PHASE I BOOK EXPLOITATION

614

~~Golodnikov, Gennadiy Vladimirovich, Nizovkina, Tat'yana Vsevolodovna  
and Ryskal'chuk, Apollinariya Terent'yevna~~

Praktikum po organicheskomu sintezu (Practical Work in Organic Synthesis) Leningrad, izd-vo Leningrad. univ-ta, 1957. 137 p.  
6,080 copies printed.

Sponsoring Agency: Leningrad. Universitet.

Ed. (title page): Dolgov, B.N., Professor; Ed. (inside book):  
Shchemeleva, Ye.V.; Tech. Ed.: Vodolagina, S.D.

PURPOSE: This manual is intended for the use of chemistry students taking a laboratory course in organic synthesis.

COVERAGE: This manual is presented as a guide to practical laboratory work in organic synthesis. It is divided into three parts. The first part is devoted to general methods and procedures of laboratory work in this field. Particular attention is given to

Card 1/E

Practical Work in Organic Synthesis

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fractional distillation, fractionating columns, steam distillation, reduced-pressure distillation, recrystallization, and the determination of melting points. Instructions are given for keeping records, safeguarding against accidents, what to do in case of fire, first aid in case of burns, glass cuts, etc. The second part describes the synthesis of organic compounds. The syntheses are divided according to types of reaction. For each type of reaction a few examples are given distinguished one from the other according to methods of separation and treatment of organic compounds produced. The third part of the manual includes supplementary syntheses which are given to laboratory students as finals. Every synthesis described in the manual has been checked and proved many times over during many years by the faculty of organic chemistry at the Leningrad State University. There are no personalities and no references.

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20-5-18/47

AUTHORS:

Dolgov, B. N. , Golodnikov, G. V. , and Golodova, K. G.

TITLE:

On the Possibility of Catalytic Dehydrogenation of Silicon-Hydrocarbons (O vozmozhnosti kataliticheskogo degidrirovaniya kremsko-levodorov)

PERIODICAL:

Doklady AN SSSR, 1957, Vol. 117, Nr 6, pp. 987 - 989 (USSR)

ABSTRACT:

There exist no references to this kind of dehydrogenation of silicon-paraffins in publications. In reference 1 it is reported that under the conditions selected there the above-mentioned reaction with tetraethylsilane did not take place. The authors succeeded in finding a catalyst (placed at their disposal by Yu. A. Gorin and S. M. Monoszon) and in determining the conditions of the dehydrogenation of a mixed tetra-alkylsilane, namely trimethylbutylsilane. The nature of the above-mentioned catalyst is not mentioned in the paper. At 550 - 575 °C 6,2 - 8,6 % yields of trimethyl-butanyl-silane, calculated on the trimethylbutylsilane sent through, were obtained (table 1). The catalyst is highly stable: neither the yields of silicon-olefin change nor is silicon deposited on the catalyst. The above-mentioned yields can still be increased by repeated passage of condensates over the catalyst, as the latter contain considerable quantities of unchanged trimethylbutylsilane.

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20-6-18/47

## On the Possibility of Catalytic Dehydrogenation of Silicon-Hydrocarbons

Beside the dehydrogenation, especially at high temperatures (575 - 600°C), some side reactions take place which are connected with the thermal decomposition of trimethylbutylsilane. Of special interest is the formation of tetramethylsilane and propylene which occurs under splitting up of the C-C bond in the butyl radical. At the same temperatures a destructive hydrogenation of the formed tetramethylsilane by hydrogen, produced in the dehydrogenation of trimethylbutylsilane takes place. Theoretically the following isomers of trimethylbutenyl-silane are possible:  $(\text{CH}_3)_3\text{SiCH}=\text{CHCH}_2\text{CH}_3$  (cis- and trans-forms) (I),  $(\text{CH}_3)_3\text{SiCH}_2\text{CH}=\text{CHCH}_3$  (cis- and trans-forms) (II), and  $(\text{CH}_3)_3\text{SiCH}_2\text{CH}_2\text{CH}=\text{CH}_2$  (III). Of these, however, only trimethyl- $\gamma$ -butenylsilane (III) is known. The authors did not succeed in isolating the silicon olefin in a pure state, as the boiling points of all products and of the initial substance are supposed to lie very close to each other. The constants of the fraction 109 - 111°C, most enriched with silicon-olefin, are in table 2 compared with the properties of the known  $\gamma$ -isomer (III) and of the initial substance. The silicon-olefin obtained by the authors apparently is the  $\gamma$ -isomer (I). The absence of the  $\beta$ -isomer (II) is confirmed by the speed of the reduction of the produced silicon-olefin. Finally the absence of the  $\beta$ -isomer is confirmed by the production of a stable dibromide

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20-6-18/47

On the Possibility of Catalytic Dehydrogenation of Silicon-Hydrocarbons

of trimethyl-butenyl-silane. A kind of short experimental part with the usual data is given which is not designated as such. There are 2 tables, and 8 references, 5 of which are Soviet.

**ASSOCIATION:**

**Leningrad** State University ineni A. A. Zhdanov  
(Leningradskiy gosudarstvennyy universitet in. A. A. Zhdanova)

**PRESENTED:** August 5, 1957, by A. V. Lopchikov, Academician

**SUBMITTED:** August 5, 1957

**AVAILABLE:** Library of Congress

Card 3/3

AUTHORS: Golodnikov, G. V., Delgov, B. N., SCV/70-26-6-21/66  
Propp, L. N.

TITLE: Catalytic Synthesis of Ketones From a Mixture of n-Butyric  
Acid and Ethyl Alcohol (Kataliticheskoye polucheniye ketonov  
iz smesi n.-maslyanoy kisloty i etilovogo spirta)

PERIODICAL: Zhurnal obshchey khimii, 1958, Vol. 26, Nr 8,  
pp. 2097 - 2099 (USSR)

ABSTRACT: The authors showed already in the previous paper (Ref 1)  
that a methyl-propylketone and besides this symmetrical  
ketones, i.e. acetone and dipropyl ketone, are formed in  
in the case of passing of a mixture of acetic acid and n-butyl  
alcohol over a Cr-Mn catalyst. The suggested reaction scheme  
is based upon a dissociation of the primarily formed ester,  
of butyl acetate into aldehydes, their aldol-condensation  
and the further transformation of aldol into methyl-propyl  
ketone. The present paper is a continuation of the experiments  
which try to synthesize mixed ketones from mixtures of acids  
and alcohols. A mixture of ethyl alcohol and n-butyric acid  
was chosen. This mixture is interesting since the assumed

Card 1/3

Catalytic Synthesis of Ketones From a Mixture of  
n-Butyric Acid and Ethyl Alcohol

SOV/77-28-8-21/66

intermediate product formed in the ketone formation, the ethyl butyrate is metameric to the butyl acetate which for its part represents an intermediate product which was formed in the ketone formation of the mixture of acetic acid n.-butyl alcohol. Thus the transformations of these esters which are in the metamerismic ratio are bound to lead to one and the same aldehydes, i.e. to the acetic- and butyric aldehyde. Therefore the formation of the same ketones, i.e. of methyl-propyl ketone, acetone, and dipropyl ketone observed in the reaction confirms to a certain extent the correctness of the suggested mechanism of the ketonization of the mixtures of acids and primary alcohols. The formation of a mixed ketone is demonstrated in the given reaction scheme, where attention must be paid to R and R' of both mixtures. The conditions and results of the experiments are given in tables 1,2,3, the properties of the ketones in table 4. Cr-Mn catalyst was used as a catalyst in this work. There are 4 tables and 4 references, which are Soviet.

Card 2/3

Asymmetric Synthesis of Ketones From a Mixture of  
Acetic Acid and Ethyl Alcohol

SCV/79-28-6-21/66

ORIGIN: Leningradskiy gosudarstvennyy universitet (Leningrad State  
University)

SUBMITTED: July 2, 1957

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AUTHORS: Dolgov, B. N. (Deceased), Goltsinkov, G. V., Gensler, I. B.TITLE: Catalytic Conversion of Tetraalkyl Silanes. III. Catalytic  
Dehydrogenation of Trimethyl Propyl-, Triethyl Butyl-,  
and Trimethyl Hexyl SilanePERIODICAL: Zhurnal obshchey khimii, 1960, Vol. 30, No. 2,  
pp. 2988-2995

TEXT: In continuation of their previous paper (Ref. 2), the authors studied the dehydrogenation of silicon hydrocarbons of the series  $(CH_3)_3SiR$ , ( $R = C_3H_7, C_4H_9, C_6H_{13}$ ). The investigations were carried out with the catalyst used in the previous investigation (Ref. 2) by Yu. A. Gorin and S. M. Monozon and placed at the authors' disposal. The tetraethyl silane passed through over this catalyst at 500-600° (volume rate 30) was not dehydrogenated: Besides unchanged tetraethyl silane, triethyl silane and diethyl silane, that are formed by the splitting off of ethylene, were detected (Table 1). The gaseous products consisted of hydrogen, saturated hydrocarbons and ethylene. The silanes of the series

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Catalytic Conversion of Tetraalkyl Silanes 3/577/60/030/001/006, 015  
 III. Catalytic Dehydrogenation of Trimethyl 8001/8064  
 Propyl-, Trimethyl Butyl-, and Trimethyl Hexyl Silane

$(\text{CH}_3)_3\text{SiR}$ , however, contain no ethyl radicals and can therefore be dehydrogenated with the above catalyst. Table 2 shows the optimum conditions of the dehydrogenation of trimethyl butyl silane, and Table 3 those of the dehydrogenation of trimethyl propyl silane (depending on the temperatures and volume rates), as well as the yields of the reaction products. The reactions carried out at temperatures above  $570^\circ\text{C}$  in the presence of a catalyst proceeded according to Scheme 2 with the bonds  $\text{C}-\text{C}$  and  $\text{C}-\text{Si}$  undergoing cleavage. Only by multiple fractional distillation, in a rectifying column, it was possible to separate the fraction boiling between  $84.0$  and  $84.5^\circ\text{C}$  that (owing to the thiocyanogen number) contained 21.1% silicon olefin. The three intensive lines of the Raman spectrum of the fractions enriched with the silicon olefin are probably caused by the isomers  $(\text{CH}_3)_3\text{SiCH}=\text{CHCH}_3$  and  $(\text{CH}_3)_3\text{SiCH}_2\text{CH}=\text{CH}_2$ . The dehydrogenation product of trimethyl propyl silane appears, according to its spectral analysis, as in the case of trimethyl butyl silane (Ref. 2), to consist for the major part of the isomer of silicon olefin that contains the double bond in

Card 2/3

Catalytic Conversion of Tetraalkyl Silanes. 3/07/60/040/001/006/015  
III. Catalytic Dehydrogenation of Trimethyl 8001/8064  
Propyl-, Trimethyl Butyl-, and Trimethyl Hexyl  
Silane

$\alpha$ -position to the silicon. A temperature of  $540^{\circ}\text{C}$  and the volume rate of 45 proved to be the optimum reaction conditions for the dehydrogenation of trimethyl hexyl silane (Table 4). Also here, the Raman spectrum shows in all cases the frequencies characteristic of the double bond. There are 8 tables and 17 references: 8 Soviet, 6 US, and 3 German.

ASSOCIATION: Leningradskiy gosudarstvennyy universitet  
(Leningrad State University)

X

SUBMITTED: October 2, 1959

Card 3/3

51879

S/079/60/030/010/020/030  
B001/B066

S 3700 4 1273, 1209

AUTHORS: Golodnikov, G. V. Dolgov, B. M. (Deceased) and  
Sedova, V. F.

TITLE: Synthesis and Properties of Trialkyl-triaryl-(p-bromo-  
phenoxy)-silanes. I. Trimethyl-, Triethyl-, and Tri-  
propyl-(p-bromo-phenoxy)-silane

PERIODICAL: Zhurnal obshchey khimii. 1960, Vol. 30, No. 10.  
pp. 3352 - 3358

TEXT: Only the first member of silicon-containing bromides of the  
general formula  $p-R_2SiOC_6H_4Br$  ( $R=CH_3$ ) has so far been described in pub-  
lications (Ref.1). The authors of the present paper intended to syn-  
thesize members of the same series with  $R = C_2H_5$ ,  $C_3H_7$ , and to study  
the possibilities of their synthesis, first of all, when using the  
Grignard reagents (obtained from the above bromides) in the synthesis  
of silicon-containing aromatic alcohols and acids. The hydrolytic sta-  
bility of the ether group  $Si-C-C_{ar}$  ( $C_{ar}$  = aromatic carbon of benzene)

Card 1/3



84679

Synthesis and Properties of Trialkyl-tri-aryl-(p-bromo-phenoxy)-silanes. I. Tri-  
methyl-, Triethyl-, and Tripropyl-(p-bromo-phenoxy)-silane

S/C79/60/C30/C10/C20/C30  
B001/B056

should further be investigated. The trialkyl (p-bromo-phenoxy)-silanes newly synthesized by the authors are stable to hydrolysis; the Si-O-C<sub>ar</sub> bond remains unchanged (Ref. 2). The secondary alcohols were synthesized according to the Scheme  $R_3SiOC_6H_4MgBr + CH_3CHO \rightarrow R_3SiOC_6H_4CH(OH)CH_3$ . Alcohols with R = CH<sub>3</sub> or C<sub>2</sub>H<sub>5</sub> however, were not obtained because of their hydrolytic instability. At R = CH<sub>3</sub>, only hexamethyl disiloxane (80% yield) and a silicon-free resin resulted. When R was C<sub>2</sub>H<sub>5</sub>, triethyl-phenoxy-silane (24%) was found in addition to hexaethyl disiloxane (38%) and a large quantity of resin. Methyl-(p-tripropyl-siloxy-phenyl) carbinol was far more stable (33%). Tripropyl-phenoxy silane (20%) and a small resin quantity were obtained as side products, which indicates a comparative hydrolytic stability of the Si-O-C<sub>ar</sub> bond in the molecule of the alcohol. Molecular weight and silicon content are in good agreement with the formula  $(C_6H_4)_3SiOC_6H_4CH(OH)CH_3$ . Further

Card 2/3

1959

Synthesis and Properties of Trialkyl-tri-  
aryl-(p-bromophenoxy) silanes. I. Tri-  
methyl-, Triethyl-, and Tripropyl (p-bromophenoxy)silane

S/073/40/313/010/020/030

EC01/1066

experiments concerning the synthesis of the alcohols show that, when propyl radicals are introduced, the hydrolytic stability of  $\text{Si-C}_{\text{ar}}$  bonds increases so much that the corresponding alcohol can be separated. Acids of the general formula  $\text{p-R}_3\text{SiC}_6\text{H}_4\text{COOH}$  could not be synthesized by means of the stabilizing propyl radicals; p-oxybenzoic acid and silicon-containing cleavage products were found instead of these acids (Table 1). The effect of the magnitude of the radical on the silicon atom upon the properties of the  $\text{Si-C}_{\text{ar}}$  bond was experimentally determined by cleavage of the Grignard reagents (Scheme 2). Table 1 illustrates the cleavage of the Grignard reagents  $\text{p-R}_3\text{SiC}_6\text{H}_4\text{MgBr}$ . There are 4 tables and 2 references, 6 Soviet, 2 US, and 1 German.

ASSOCIATION: Leningradskiy gosudarstvennyy universitet (Leningrad State University)

SUBMITTED: December 2, 1959

Card 3/3

EL385

5 3700 2209, 1271, 1153

S/073/60/050/010/028/030  
300/3066

AUTHORS: Golodnikov, G. V. and Repinskaya, I. B.  
TITLE: Catalytic Dehydrocyclization of Trimethyl hexyl Silane  
PERIODICAL: Zhurnal obshchey khimii, 1960, Vol. 30, No. 10,  
pp. 3501 - 3502

TEXT: In order to find out whether dehydrocyclization of trimethyl-  
hexyl silane is possible, the authors allowed it to pass over a chromium  
catalyst at 530-590°C (Ref.1). Under these conditions, dehydrocycliza-  
tion gives trimethyl-phenyl silane:

$(CH_3)_3SiC_6H_{13} \rightarrow 4H_2 + (CH_3)_3SiC_6H_5$  The aromatization of silicon paraf-  
fins is a reaction hitherto unknown in organosilicon chemistry. An un-  
saturated hydrocarbon appears as intermediate, whose content in the con-  
densates varied between 7.5 and 21%, according to the temperature (de-  
termined by the thiocyanation method). Trimethyl silane, tetramethyl  
silane, pentene, hexene and benzene were the side products. The con-  
densates repeatedly underwent fractional distillation. The separated

Card 1/2

ELSES

Catalytic Dehydrocyclization of  
Trimethyl-hexyl Silane

S/070/60/C30/010/028/030  
B001/B066

fraction with a boiling point between 163 and 165° contained trimethyl-phenyl silane in addition to unchanged trimethyl hexyl silane. Ref. 2 gives the data published on trimethyl-hexyl silane, and Ref. 3 those on trimethyl-phenyl silane. The presence of trimethyl-phenyl silane in the fraction boiling between 163 and 165° is confirmed by its refractive index and spectroscopic data. The infrared spectrum of the fraction shows an absorption maximum which is characteristic of the phenyl ring which is bound to the silicon (Ref.4) This study is being continued to find stronger catalysts for the dehydrocyclization of silicon paraffins. There are 4 references: 1 Soviet, 1 Japanese, and 2 US. ✓

ASSOCIATION: Leningradskiy gosudarstvennyy universitet (Leningrad State University)

SUBMITTED: May 26, 1960

Card 2/2

S 015761 031 011 001 001  
0118 0101

AUTHORS

Golodnikov, G. V. and Abramovskiy, A. I.

TITLE

Tetramyl-, trihexyl- and triphenyl-(n-bromophenoxy)silanes

PERIODICAL:

Zhurnal obshchey khimii, 36, no. 11, 1966, 3352-3353, 13-

TEXT:

The authors studied the synthesis and properties of three little known silicoorganic bromides and two new silicoorganic alcohols: tetramyl-(n-bromophenoxy)silane (I); trihexyl-(n-bromophenoxy)silane (II); triphenyl-(n-bromophenoxy)silane (III); methyl-(n-bromophenoxyphenyl)carbinol - a  $(C_5H_4)_3SiOC_6H_4CH(OH)CH_3$  (IV) and methyl-

(n-trihexylsiloxyphenyl)carbinol - a  $(C_6H_{11})_3SiOC_6H_4CH(OH)CH_3$  (V). Previous research by G. V. Golodnikov, B. K. Dolgov, A. F. Zolotarev (Zh. obshch. khimii, 36, 3352, 1966) on the properties and reactions of bromides of the type  $aR_3SiOC_6H_4Br$  disclosed the possibility of their

Card 1/3

Triamyl-, trihexyl- and

conversion into secondary silicoorganic alcohols by means of organohydro- and magnesioorganic compounds. The method of the catalytic reduction and condensation of trialkyl- and triaryl silanes with a triphenylborane- $ZnCl_2$  or  $ZnEt_2$  catalyst, given in R. N. Depp, V. I. Anisimov, N. F. Khazanova (Ref. 12, AN SSSR, Otdel. Khim. i Mekh. 1981) was used to prepare I, II, and III. For syntheses of IV, a mixture of Mg, abs. ether, I, and ethyl bromide was placed in a flask, after which the solution was successively cooled, reacted with ethyl bromide, abs. ether, boiled for a further 50 min, and filtered with pressure. The subsequent vacuum distillation of the residue yielded three fractions: 175-205° (containing phenylsilane, 10%), 205-225° (contaminated with some unreacted triphenylborane, 10%), and 225-245° (II). II was converted into the desired alcohol by means of the procedure of fraction boiling at 80-120° containing a propane gas stream, phenylsilane, while the 205-225° fraction contained triphenylborane and triethylsilane, and the 225-245° fraction contained triphenylborane and triethylsilane. The authors considered by the authors to be of interest for the synthesis of silanes.

Card 20

376797617031701370137013  
D22870405

Triamyl-, trihexyl-, and

Si-O-C bond in the molecules of IV and V, which may be due to the screening effect of the amyl and hexyl radicals linked with the Si atom. IV and V react with Na to give  $H_2$ , and their IR (K) spectra contain absorption maxima at  $3400\text{ cm}^{-1}$  and  $3380\text{ cm}^{-1}$  respectively. There are 2 tables and 4 references: 2 Soviet-bloc and 2 non-Soviet bloc.

ASSOCIATION: Leningradskiy gosudarstvennyy universitet (Leningrad State University)

SUBMITTED: December 6, 1960

Card 3/3

S/079/61/031/011/014/015  
D228/D305

5 3700  
AUTHORS:

Golodnikov, G. V., and Koroleva, G. N.

TITLE:

Catalytic dehydrogenation of trimethylethylsilane

PERIODICAL:

Zhurnal obshchey khimii, v. 31, no. 11, 1961, 3738-3740

TEXT:

The authors studied the catalytic dehydrogenation of trimethylethylsilane—the first member of a group of mixed silanes with the formula  $\text{Me}_3\text{SiR}$ . B. N. Dolgov, G. V. Golodnikov, and I. B. Gensler (Ref. 1: Zh. obshch. khimii, 30, 2988, 1960), whose experimental procedure was followed in this work, also examined the dehydrogenation of other mixed silanes— $\text{Me}_3\text{SiPr}$ ,  $\text{Me}_3\text{SiHx}$ —and showed that tetraethylsilane, on the contrary, does not undergo dehydrogenation under the chosen conditions. The authors' data indicate that the reaction proceeds best at 590 – 600°, some 20 – 30° higher than is the case with silanes containing propyl, butyl, and hexyl radicals. Raising the temperature to 620° promotes the development of side-reactions—when trimethylsilane (I), trimethylvinylsilane (II),

Card 1/2



AUTHORS: Golodnikov, G. V. and Khalutina, A. A.

S:079/62/032/007/003/007  
1032/1232

TITLE: Catalytic dehydrogenation of  $\gamma$ -trimethyl silyl-propyl alcohol

PERIODICAL: Zhurnal obshchei khimii, v. 32, no. 7, 1962, 2302-2305

TEXT: The aim of this study was to work out a general method for the preparation of silicon-containing aldehydes by way of dehydrogenation of the corresponding primary alcohols. The best yield of  $\beta$ -trimethyl-silyl-propionic aldehyde was obtained by dehydrogenation of the corresponding alcohol over a catalyst designated "violet copper" (reduced copper) at 300 °C. The yield amounted to 76.7% of the alcohol that reacted, or to 37.8% of the total amount of alcohol. 45% of the alcohol did not react. Side reactions took place to a small extent. Dehydration gave trimethyl-allyl-silane (3.5%). Hexamethyl-disiloxane (2.3%) and propylene were also formed. There is 1 table. The English-language references read: 1) K. Frisch, and P. Shroff, J. Am. Chem. Soc., 75, 1249 (1953); Ch. A., 47, 9055 (1953). 3) C. Burkhard and D. Hurd, J. Org. Ch., 17, 1107 (1952). 4) C. Brannen Ch. A. 48, 624 (1954). 5) J. Speier, J. Webster, G. and Barnes, J. Am. Chem. Soc., 79, 574 (1957). 9) L. Sommer, R. Van Streen and F. Whitmore, J. Am. Chem. Soc., 71, 3056 (1949). 6) Beilstein, 162, 163, 164, 165, 166, 167, 168, 169, 170, 171, 172, 173, 174, 175, 176, 177, 178, 179, 180, 181, 182, 183, 184, 185, 186, 187, 188, 189, 190, 191, 192, 193, 194, 195, 196, 197, 198, 199, 200.

ASSOCIATION: Leningradskii gosudarstvennyi universitet

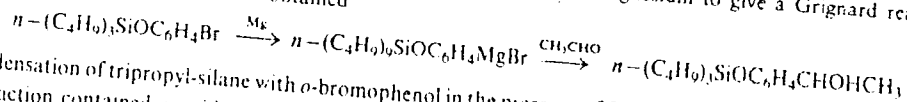
AUTHORS: Golodnikov, G. V. and Sycheva T. P.

S:079/62/032/007/004/007  
I032/I232

TITLE: Synthesis and properties of tributyl-(*p*-bromophenyl)-silane

PERIODICAL: Zhurnal obshchei khimii, v. 32, no. 7, 1962, 2305-2307

TEXT: Tributyl-(*p*-bromophenoxy)-silane, a compound not described in the literature hitherto, was synthesized by catalytic condensation (involving dehydrogenation) of tributyl-silane with *p*-bromophenol, in the presence of SnCl<sub>2</sub>. Tributyl-(*p*-bromophenoxy)-silane reacted with magnesium to give a Grignard reagent, from which a carbinol could be obtained



Condensation of tripropyl-silane with *o*-bromophenol in the presence of SnCl<sub>2</sub> was attempted, but the products of reaction contained considerable amounts of hexapropyl-disiloxane, *o*-bromophenol, as well as 57.9% of the calculated amount of hydrogen. The tripropyl-(*o*-bromophenoxy)-silane is considered to have undergone hydrolytic dissociation.

ASSOCIATION: Leningradskii gosudarstvennyi universitet (Leningrad State University)

SUBMITTED: July 5, 1961

Card 1/1

GOLODNIKOV, G.V.; D'YAKONOV, I.A.; REPINSKAYA, I.B.; FOMINA, O.S.

Copper sulfate catalyzed reaction of diazoacetic ester with  
3-trimethylsilyl-1-propene and 4-trimethylsilyl-1-butene.  
Zhur.ob.khim. 33 no.7:2422-2423 J1 '63. (MIRA 16:3)

1. Leningradskiy gosudarstvennyy universitet.  
(Silicon organic compounds) (Acetic acid)

GOLDBERGOV, G.V.; SHARSHALINA, V.V.

Catalytic dehydrogenation of  $\gamma$ -trialkylsilylpropyl alcohols.  
Part 2. Zhur.ob.khim. 33 no.10:3262-3264 O 1983.(MIRA 16:11)

1. Leningradskiy gosudarstvennyy universitet.

GOLODNIKOV, G.V.; MARKIN, G.A.

Synthesis and properties of trialkyl(p-bromophenoxy)silanes. Part 4.  
Zhur.ob.khim. 33 no.10:3265-3266 O '63. (MIRA 16:11)

1. Leningradskiy gosudarstvennyy universitet.

D'YAKONOV, I.A.; GOLODNIKOV, G.V.; REPINSKAYA, I.B.; FOMINA, O.S.

Reactions of diphenylmethylene and carbethoxycarbene with  
alkenylsilanes. Zhur.ob.khim. 33 no.10:3438-3439 O '63.  
(MIRA 16:11)

1. Leningradskiy gosudarstvennyy universitet.



D'YAKONOV, I.A.; GOLODNIKOV, G.V.; REPINSKAYA, I.B.

Reactions of aliphatic diazo compounds with unsaturated compounds.  
Part 25: Reaction of diphenyldiazomethane with silicon olefins.  
Zhur.org.khim. 1 no.2:220-225 F '65.

(MIRA 18:4)

1. Leningradskiy gosudarstvennyy universitet.



D'YAKONOV, I.A.; SEPINSKAYA, I.B.; GOLCDNIKOV, G.V.

Trimethylsilylcarbene, a new methylene radical. *Dokl. Akad. Nauk SSSR*  
35 no.1:199 Ja '65. [PTA 1814]

1. Leningradskiy gosudarstvennyy universitet.

D'YAKOV, I.A.; GOLODNIKOV, G.V.; PRUDENSKIY, L.B.

Reactions of aliphatic diazo compounds with substituted alkenes.  
Part 25: Reaction of ethyl ester of diazoacetic acid with  
trimethylvinyl-, trimethylallyl-, and trimethyl- $\alpha$ -phenylallenes.  
Zhur.b.khim. 35 no.12:2181-2185, 1965. (1965:10:1)

I. Leningradskiy gosudarstvennyy universitet. Laboratoriya khimii  
i, 1964.

L 45894-60 ENT(m)/ENT(j) WH/RM

ACC NR: AP6026430

(A)

SOURCE CODE: UR/0079/56/036/005/0949/0949

AUTHOR: D'yakonov, I. A.; Repinskaya, I. B.; Golodnikov, G. V.

ORG: Leningrad State University (Leningradskiy gosudarstvennyy universitet)

TITLE: Relative rate of addition of diphenylmethylene to trimethylvinyl-, trimethylallyl- and trimethyl- $\gamma$ -butenylsilanes and 1-heptene

SOURCE: Zhurnal obshchey khimii, v. 36, no. 5, 1966, 49

TOPIC TAGS: silane, heptene, organosilicon compound

ABSTRACT: Diphenylmethylene (DM), obtained by thermal decomposition of diphenyldiazomethane (DD), adds to the silicoolefins (SO) trimethylvinyl-, trimethylallyl- and trimethyl- $\gamma$ -butenylsilanes, forming silicon-containing cyclopropane (CD). An estimate of the relative activity of SO in their reaction with DD was made, based on the use of the method of competing reactions, and on the determination of the relative rate constant of the reaction ( $K_{rel}$ ) of each of the SO. The reference standard was 1-heptene, whose rate constant of the reaction with DD was taken as unity. The  $K_{rel}$  values obtained were:  $(CH_3)_3SiCH=CH_2$ , 46.3;  $(CH_3)_3SiCH_2CH=CH_2$ , 0.93;  $(CH_3)_3SiCH_2CH_2CH=CH_2$ , 0.91. These values indicate that the steric effect of the trimethylsilyl group in trimethylvinylsilane has no appreciable effect on the rate of addition of DM to the double bond of this silane. The product of the addition of DM to 1-heptene (16% yield).

Card 1/2

UDC: 547.6+547.245

L 45894-66

ACC NR: AP6026430

1-amyl-2,2-diphenylcyclopropane, is described for the first time.

SUB CODE: 07/ SUBM DATE: 12Oct65/ ORIG REF: 003/ OTH REF: 004

Card 2/2 *LC*

65

GOLODNIKOV, N.A., master

Electric tachometer. *Energetik* 8 no.8:27-28 Ag '60. (MIRA 13:10)  
(Tachometer)

GOLODNOV, A.; GUSEVA, N., red.; MAGIBIN, P., tekhn. red.

[Sheep raising as a basic source of meat production]  
Ovtsevodstvo - osnovnoi istochnik proizvodstva miasa.  
Alma-Ata, Kazsel'khozgiz, 1962. 26 nos. in 1 v. 22 p.  
(MIRA 17:1)

SALYUKOV, P.A., kand. biol. nauk, VERNIGOR, V.A., kand. sel'khoz. nauk; KORMANOVSKAYA, M.A., kand. sel'khoz. nauk; GOLOLNOV, A.V.; SKOROBOGATOV, Yu.A., mladshiy nauchnyy sotr.; MALLITSKIY, V.A., kand. sel'khoz. nauk; CHAPCHIN, B.V., kand. sel'khoz. nauk; PONOMAREV, P.P., kand. tekhn. nauk; BARVINTSEV, Yu.N., doktor sel'khoz. nauk; NECHAYEV, I.N., mlad. nauchnyy sotr.; POZDNYAKOV, P.M., kand. biol. nauk; KOVIN'KO, D.A., kand. biol. nauk; BAIANINA, O.V., kand. sel'khoz. nauk; MOISEYEV, K.V., kand. sel'khoz. nauk; ROMANOV, P.F., kand. veter. nauk; PAI'GOV, A.A., kand. veter. nauk; ANAN'YEV, P.K., kand. veter. nauk; VASIL'YEV, B.M., kand. sel'khoz. nauk; ABDULLIN, V.A., kand. ekon. nauk; GALIAKBEROV, N., laureat Gos. premii, kand. sel'khoz. nauk, red.; GUSEVA, N., red.; NAGIBIN, P., tekhn. red.

[Reference book for zootechnicians] Spravochnik zootekhnika. Pod red. N. Galiauberova. Alma-Ata, Kazsel'khozgiz, 1963. 492 p. (MIRA 16:5)  
(Kazakhstan--Stock and stockbreeding)



*Journal of Management Education*, 20(6), 709-728.

"Instruction, training of" (as per sense "class, ill. application of the system," per May 11, 1914) (see correspondence to graduate school, Minister of Higher Education 1914)

Dissertations presented for release on 1-21-67 by the Library of Congress in Moscow, April - 1967.

Ch. J. W. L., 6 May '66

AID P - 3435

Subject : USSR/Electricity  
Card 1/2 Pub. 27 - 2/32  
Author : Golodnov, M. N., Kand. of Tech. Sci., Dotsent,  
Rostov-on-the-Don  
Title : Permissible load in starting synchronous motors with  
directly-connected exciter  
Periodical : Elektrichestvo, 10, 7-10, 0 1955  
Abstract : Current circulars of the Ministry of Electric Power  
Stations No. 7/E 1952 and No. E-5/54 1954 permit the  
starting without preliminary tests of synchronous  
motors with directly connected exciter and with  
resistance moment not exceeding 0.4 of the nominal.  
The author presents a simple engineering method of  
approximate calculations to determine the possibility  
of applying direct starting of synchronous salient-  
pole motors. He presents results of calculating the  
characteristics of motors of the SM-300-750 217-kw,

Elektrichestvo, 10, 7-10, 0 1955

AID P - 3435

Card 2/2      Pub. 27 - 2/32

6-kv, and SM-160-500 113-kw, 380-v types. The author suggests experimenting with his method to ascertain the possibility of starting motors with a resistance moment above 0.4 of the nominal. Analytical findings of the author have been verified experimentally not only by himself, but also by several laboratories (Central Scientific Research Institute of the Ministry of Electric Power Stations and others). Two diagrams, 2 Soviet references (1950, 1952).

Institution : None

Submitted : N 29, 1954

AUTHORS: Golodnov, L. N., Docent, Candidate of Technical Sciences (Moscow) 105-58-4-22/37

TITLE: Experimental Determination of the Parameters of an Equivalent Generator in Linear A. C. Circuit  
(Opytnoye opredeleniye parametrov ekvivalentnogo generatora v lineynykh tsepyakh peremennogo toka)

PERIODICAL: Elektrichestvo, 1958, Nr 4, pp. 77-78 (USSR)

ABSTRACT: The determination of the electromotive force of an equivalent generator can be determined without difficulty from the idle motion of a dipole. It is useful to determine the total resistance of the equivalent generator in trying to feed the dipole from a foreign source. This should be done by measuring it at the terminals of the dipole as input resistance while all energy sources feeding it are disconnected and replaced by circuit sections with the same internal resistances. Sometimes, however, such a disconnection is not possible or is not desired. In such a case the parameters  $r_H$  and  $x_H$  can be determined by two experiments applying different loads to the effective dipole.

Card 1,3

Experimental Determination of the Parameters of an  
Generator in Linear A. C. Circuit

105-58-4-22/57

After the idle motion voltage  $U_0 = E_H$  was determined the load resistance  $Z_1 = r_1 + jx_1$  is connected to the dipole terminals. Then

$$I_1 = \frac{U_0}{\sqrt{(r_H + r_1)^2 + (x_H + x_1)^2}} \quad (1)$$

$I_1$  is determined according to the amperemeter indication and  $r_1$  and  $x_1$ , if unknown, by the indications of amperemeter, voltmeter and wattmeter. The resistance  $Z_1$  is replaced by  $Z_2 = r_2 + jx_2$  and the equation (2) is obtained. Both equations (1) and (2) are solved in common and the resistance  $r_H$  and  $x_H$  are determined. In practice mostly only effective resistances  $r_1$  and  $r_2$  can be connected. Thus

Page 2/3

Experimental Determination of the Parameters of an  
Equivalent Generator in Linear A. C. Circuit

105-58-4-2/37

the scheme of the experiment becomes more simple (the  
wattmeter is no longer needed). The method was tried in  
practice and the results were good.

AVAILABLE: Library of Congress

1. Generators-Parameter determination-Theory 2. Linear circuits

Card 3/3

AUTHOR: Golodnov, M. N. Candidate of Technical Sciences 1c5-58-8-27/77  
(Rostov-na-Donu = Rostov on Don)

TITLE: The Determination of the Synchronous Reactance of Three-Phase-Current Generators With Salient Poles and Permanent Magnets in an Experimental Way (Opredeleniye opytnym putem sinkhronnogo reaktivnogo soprotivleniya trekhfaznykh yavopolysnykh generatorov s postoyannymi magnitami)

PERIODICAL: Elektrichestvo, 1958, Nr 6, pp. 91-91 (USSR)

ABSTRACT: The method of the small slip which is employed for the determination of the synchronous reactances  $x_d$  and  $x_q$  of motors with salient poles cannot be used for motors with permanent magnets. In this case it is expedient to employ an indirect method for the determination of  $x_d$  and  $x_q$  by means of a test with two loads. This method is described here. During idling of the generator the linear voltages  $U_{ab}$ ,  $U_{bc}$ ,  $U_{ca}$ , are measured and the no-load e.m.f.  $E_0$  is determined ... equation(1). An inductive symmetrical three-phase load is connected with the generator, where the load possesses a low effective resistance. ( $\cos \phi = 0.1-0.2$ ). Then the amperages and the linear voltages of the generator are measured. Corresponding to the

Card 1/2

The Determination of the Synchronous Reactance of Three-Phase-Current Generators With Salient Poles and Permanent Magnets in an Experimental Way 105-55-6-27/81

vector diagram the formula (2) is then written down and from it the formula (3) for  $x_d$  is obtained. As the angle  $\varphi$  is near to  $90^\circ$  and differs little from  $\Psi$ , (3) can be written down in the form of (4). For the determination of  $x_q$  a symmetrical load of a capacity nature is produced in a way that current  $I$  according to its phase coincides with the e.m.f.  $E_0$ .  $x_q$  is determined from equation (6). There are 2 figures.

1. Generators--Performance 2. Generators--Analysis 3. Magnets  
--Electrical effects 4. Mathematics

Card 2/2



SOV/144-58-7-6/15

AUTHOR: Golodnov, Mikhail Nikolayevich, Candidate of Technical Sciences, Docent

TITLE: Equations of the Additional Current and Torque that Characterise the Influence of the Exciter e.m.f. on Asynchronous Operation of a Synchronous Machine (Uravneniya dopolnitel'nogo toka i dopolnitel'nogo momenta, kharakterizuyushchiye vliyaniye e.d.s. vzbuditelya na asinkhronnyuyu rabotu sinkhronnoy mashiny)

PERIODICAL: Izvestiya vysshikh uchebnykh zavedeniy, Elektromekhanika, 1958, Nr 7, pp 57-66 (USSR)

ABSTRACT: The meaning of the term 'additional currents' is first defined as follows: If the equations of a synchronous machine are expressed in operator form the stator and rotor currents may be divided into two components. The first of these are currents due to the system voltage applied to the stator circuit which do not depend on the exciter e.m.f. These currents can be observed under asynchronous (or synchronous) conditions in the absence of excitation and they are termed 'asynchronous currents'. Secondly, the currents due to the presence of the exciter e.m.f. in the closed field circuit. In

Card 1/6

SOV/144-58-7-6/15

Equations of the Additional Current and Torque that Characterise the Influence of the Exciter e.m.f. on Asynchronous Operation of a Synchronous Machine

the presence of excitation these currents are superposed on the 'asynchronous currents' and are termed 'additional' currents. If the effects of saturation and hysteresis are neglected the effect of the asynchronous and additional currents may be considered separately. The equations for the additional current and the additional torque together with the equations for the asynchronous current and torque may be used to analyse the operation of synchronous machines running under asynchronous conditions with the exciter connected. In particular these equations have been useful in analysis of the following conditions: starting of synchronous motors with the exciter connected; self-synchronisation of generators; asynchronous operation when the field circuit is open; and some cases of loss of synchronism of a motor. The operator equations for the additional currents are given in expressions (10) - (16). Expressions are then derived for the additional current when the field

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current is constant, which is the case for all practical purposes if the exciter is left connected and the field rheostat is not suddenly altered during the process. A number of practical conclusions are derived from the equations. Test results of the additional current and short circuit current as functions of the field current are given in Fig 2 for nine different motors; the test results are stated to confirm equations (31) - (33) and the conclusions drawn from them. The additional current was determined from oscillograms as shown in Fig 1. The characteristics obtained were compared with short circuit characteristics and were usually found to coincide, but in some cases the additional current characteristic lies somewhat lower than the short circuit characteristic, though as will be seen from Fig 2 the difference is not great. The torque expression (3) contains some terms which depend on the angle  $\delta$  and some which do not. Accordingly the additional torque can be resolved into two components, a mean value given by expression (34)

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and a periodic component given by expression (35). By further analysis it is shown that the periodic component of the additional torque does not depend on the slip and varies sinusoidally with the slip frequency. An oscillogram of this torque as a function of  $\delta$  with constant slip is given in Fig 3. Expression (39) is derived for the mean value of the additional torque. It will be seen that in general the mean additional torque is small and it is important only in a few special cases. One case in which the additional torque is important is when on starting a synchronous motor against a load the machine fails to run up to synchronous speed because of troughs in the torque/speed curve. It is shown that in some cases if the field current is increased when this happens the torque may become less because of the retarding effect of the mean additional torque, and then it is necessary either to reduce the mechanical load or to introduce active resistance into the field circuit so that the machine can run up to

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speed. Test data showing results of increasing the field current when machines fail to run up to speed are given in Table 1, which also includes a few torque values. The assumptions made in the article limit the field of application of eqs (29), (33), (38) and (39) to conditions where the slip is much less than 1, in large modern machines of fairly large flywheel effect with low active stator resistance and a massive rotor. The limitations imposed in the derivation of formulae (38) and (39) render them unsuitable for use at half synchronous speed. Thus Eqs (29) and (38) and conclusions drawn from them can be used for the relatively common case of a machine operating under synchronous conditions with low active resistance and slip less than 0.5. Under other conditions and particularly at low speeds there may be other components of additional current and torque which may be of

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considerable importance. In such cases it is best to  
use Eqs (20) - (24) or graphical methods which are not  
considered here.

There are 6 figures, 1 table and 9 Soviet references.

ASSOCIATION: Kafedra elektrotekhniki Rostovskogo instituta  
sel'skokhozyaystvennogo mashinostroyeniya (Chair of  
Electrical Engineering, Rostov Institute of  
Agricultural Machine Building)

SUBMITTED: January 31, 1956

Card 6/6



GOLODNOV, Yu.M.

New method for determining the angle between electromotive force  
vectors and the voltage on the rings of single-phase converter.  
Sbor.rats.predl.vnedr.v proizvod. no.5:50-52 '60. (MIRA 14:8)

1. "Yuvenergochermet".  
(Electric current converters)



GOLODNOV, Yu.M., inzh.

Some characteristics of a rectifier cascade. Vest.elektrom.  
31 no.1:21-24 Ja '60. (MIRA 13:5)  
(Electric motors, Induction) (Electric driving)

9.2200 (1001, 1482)

32194  
S/196/61/000/010/022/037  
E194/E155

AUTHOR: Golodnov, Yu. M.

TITLE: A direct current amperes-squared-hour meter

PERIODICAL: Referativnyy zhurnal, Elektrotehnika i energetika,  
no. 10, 1961, 1 abstract LOK 6 (Vestn elektroprom-  
sti, no. 4 1961, 56-60)

TEXT: "Yuvenergochermet" has developed and made a direct  
current amperes-squared-hour meter, using compact amplidyne.  
The electrical circuit and winding data are given, and the  
construction and operating principles are described. The meter  
reading is completely independent of the direction of current in  
the amplidyne control winding. Any single-phase induction meter  
for 127 V, 5 A can be used in the circuit. With a rated control  
winding current of 2.5 A the meter constant is 6.52 A<sup>2</sup>h/kWh. The  
equipment fits into a carrying case and is easily transported. ✓  
A variant is described in which the secondary winding of the  
amplidyne carries 5 A. This set consumes less power, so that a  
150 VA stabiliser can be used. With a rated control winding  
current of 1.5 A the meter constant is 0.965 A<sup>2</sup>h/kWh.  
Card 1/2

32191

A direct current amperes-squared.

S/196/61/000/010/022/037

E194/E155

An important advantage of the meter is its ability to operate with a shunt for 33 - 45 mV from which it draws a very small current. Shunts for 75 and 100 mV are used in conjunction with additional resistances. The meter can be used to measure r.m.s currents in reversing circuits having peak loads with an accuracy of 2.5%. The meter can be used in circuits of 100 A and more 5 figures.

X

[Abstractor's note: Complete translation]

Card 2/2

GOLODNOV, u.m., inzh.

D.C. ampere square hour meter. Vest. elektropram. 32 no.4:56-60  
Ap 1961. (MIRA 15:5)

(Electric meters)

(Electric machinery--Measurements)

SATPAIYVA, T. I.; GOLUBEVA, N. B.

Isopykto in the Dzheskizgan mines. Izv. Akad. Nauk Ser. geol.  
1977 10:112-117, (MLRA 10:8)  
(Ozhonovskan-Soviet press)

GOLODNOVA, O.S., inzh.

Review of L. A. Mirenburg's book "Repair of turbogenerator rotors  
under stationary conditions." Elek. sta. 31 no. 12:88 D '60.

(MIRA 14:5)

(Turbogenerators--Maintenance and repair)  
(Mirenburg, L.A.)

GURVICH, V.S., inzh.; GOLODNOVA, O.S., inzh.

Rotor bands from aluminum alloys. Elek.sta. 32 no.4:94-95 Ap  
'61. (MIRA 14:7)  
(Turbogenerators)

GOLODNOVA, O.S., inzh.; DEGIL', G.S., inzh.; PANCHENKO, A.U., inzh.;  
TUROS, A.E., inzh.; MESHKOV, V.K., inzh.

Concerning the seals of hydrogen cooled turbogenerators. Elek.  
sta. 33 no.8:60-68 Ag '62. (MIRA 15:8)

1. Rostovenergo (for Golodnova). 2. Glavnoye upravleniye  
energeticheskogo khozyaystva Donetskogo basseyna (for Degil',  
Panchenko, Turos). 3. Moskovskoye rayonnoye upravleniye  
energeticheskogo khozyaystva Glavtsentroenergo Ministerstva  
elektrostantsiy SSSR (for Meshkov).  
(Turbogenerators)



GOLODNOVA, O.S., inzh.

Choice of the dimensions of shaft grease seals. Vest. elektroprom.  
33 no.8:72-73 Ag '62 . (MIRA 15:7)  
(Turbogenerators--Equipment and supplies)

GOLODNOWA, O.S., inzh.

Expenditure of hydrogen in the cooling of turbogenerators. Elek. sta.  
36 no.6:50-53 Ja '65. (MIRA 1887)

GOLODNYAK, G. S.

Automation of production processes at the Semipalatinsk cement  
plant. TSement 27 no.5 24-27 S-O '61. (MIRA 14 12)

1. Semipalatinskiy tsementnyy zavod.  
(Semipala'tinsk Cement plants)

GOLODNYAK, G.S.

Automatic sprinkling of the shells of cement mills. Tsement 28  
no.1:21 Ja-F '62. (MIRA 16:5)

1. Bezmeinskiy tsementnyy zavod.  
(Milling machinery)

GOLODNYAK, N.

Encourage the activities of key personnel. Voen.znan. 35 no.4:  
11-12 Ap '59. (MIRA 12:7)

1. Predsedatel' Kiliyskogo rayonnogo Komiteta Vsesoyuznogo  
dobrovol'nogo obshchestva sodeystviya armii, aviatsii i flotu.  
(Military education)

USSR / Farm Animals. Small Horned Sheep

Abstract: Ref Zhur-Biol., No 5, 1958, 21471

Author : Olenin F. S., Gritsay S. P., Golodnyy I. F.

Inst :

Title : A New Technique of Feeding Lambs and Its Effectiveness (Novaya tekhnika kormleniya yagnyat i ego ekonomicheskaya effektivnost')

Orig Pub: S. kh. Povolzh'ya, 1957, No 6, 67-69

Abstract: Laboratory experiments were carried out in order to test "rhythmic" feeding onweanling lambs (the ration of one 6-day period was increased by 20% and that of the other 6-day period was diminished by 20%). It was found that the average daily weight increase of test animals was 32%, shearing yield 400 g. more, and the feeding per 1 kg. of weight increase was 2.42 feed units less than in the control group.

Card 1/2

AUTHOR: Golodnyy, Ts. 25-11-24/23  
TITLE: Explorer of Arctic Regions (Issledovatel' Arktiki)  
PERIODICAL: Nauka i Zhizn', 1957, # 11, pp 57-58 (USSR)  
ABSTRACT: Petr Filimonovich Shvetsov, doctor of geological-mineralogical sciences, member-correspondent of the USSR Academy of Sciences, winner of the Stalin prize, succeeded the Soviet geologist Vladimir A. Obrushev as head of the Institute for the Study of Congelation. Soviet scientists, such as Gumgin, Tsytovich, Tolstikhin, Kudryavtsev, Baranov, Soltykov, prepared the ground for this new branch of science. Shvetsov was among the first scientists to recommend the exploration of the arctic regions. He established the first Soviet station for the study of congelation on the Chukotsk peninsula. His first scientific publication dealt with the research of the Pinkegneysk thermal spring, and he provided the possibility to take advantage of these valuable resources. The results of his second expedition to the Chukotsk peninsula were summed up in his second work "Eternal Congelation and Technical Geological Conditions in the Anadyrsk Region". The material for his latest publication

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Explorer of Arctic Regions

25-11-24/28

"Subterranean Waters of the Verkhoyansk-Kolyma Folded Region and Peculiarities of Their Occurrence, Connected with Eternal Congelation", was collected during a two-year research expedition to Yakutiya.

There are three photographs and one sketch.

AVAILABLE: Library of Congress

Card 2/2



GOLODNYY, TS .

~~Production line~~ Production line system in building apartment houses. Nauka  
i zhizn' 24 no.3:21-24 Mr '57. (MLRA 10:5)  
(Apartment houses)

AUTHOR: Golodnyy, Ts.

86-1-21044

TITLE: An Experimental Block (Kvartal. eksperimentalny)

PERIODICAL: Nauka i Zhizn', 1986, # 1, pp 68-74 (USSR)

ABSTRACT: This article deals with the construction of the first Soviet experimental apartment buildings in the Novyye Cheremuski area of Moscow. The building ground consists of two parts: the housing area and the social center. There are 13 four-storied and 5 eight-storied apartment houses under construction. The social center consists of a school, a kindergarten, a public nursery, repair services, shops, a public dining-room and parking lots.  
There are seven photographs and one illustration.

AVAILABLE: Library of Congress

Card 1/1

GOLODNYI, TS.

At the anniversary exhibition. Rabotnitsa 36 no.1:24 Ja '58.  
(MIRA 11:2)  
(Moscow--Art--Exhibitions)

GOLODNYI, TS.

A book arrived at the plant. Rabotnitsa 37 no.10:27 0 '59.  
(MIRA 13:2)

1. Kombinat "Trekhgornaya manufaktura," Moskva.  
(Moscow--Booksellers and bookselling)